

Please amend specification passage at page 6, lines ¹⁻¹⁹~~3-21~~ as follows:

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Suitable values of R^1 , R^2 , R^3 , R^4 , R^5 and p are as follows. Such values may be used where appropriate with any of the definitions, claims or embodiments defined hereinbefore or hereinafter.

R^1 is fluoro, chloro, cyano, methyl, ethyl, methoxy or ethoxy.

p is 0.

p is 1.

p is 2.

R^2 is $R^6-NH-R^6-NH_2$ wherein R^6 is C_{1-4} alkyl, C_{2-4} alkenyl, C_{3-6} cycloalkyl, C_{3-6} cycloalkyl C_{1-3} alkyl or (heterocyclic group) C_{1-3} alkyl; and wherein R^6 may be optionally substituted on carbon by one methoxy, ethoxy or trifluoromethyl.

R^2 is $R^6-NH-R^6-NH_2$ wherein R^6 is C_{1-4} alkyl, C_{2-4} alkenyl, C_{3-6} cycloalkyl, C_{3-6} cycloalkyl C_{1-3} alkyl or (heterocyclic group) C_{1-3} alkyl; and wherein R^6 may be optionally substituted on carbon by one methyl, methoxy, ethoxy or trifluoromethyl.

R^2 is $R^6-NH-R^6-NH_2$ wherein R^6 is methyl, ethyl, propyl, *t*-butyl, allyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, tetrahydrofur-2-ylmethyl or pyrid-2-ylmethyl; and wherein R^6 may be optionally substituted on carbon by one methoxy, ethoxy or trifluoromethyl.

R^2 is $R^6-NH-R^6-NH_2$ wherein R^6 is methyl, ethyl, propyl, *t*-butyl, allyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, tetrahydrofur-2-ylmethyl or pyrid-2-ylmethyl; and wherein R^6 may be optionally substituted on carbon by one methyl, methoxy, ethoxy or trifluoromethyl.

Please amend specification passage at page 7, lines ¹³⁻²⁸~~15-30~~ as follows:

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Therefore in another aspect of the invention, there is provided a compound of formula (I) (as depicted above) wherein:

p is 0;

R^2 is $R^6-NH-R^6-NH_2$ wherein R^6 is C_{1-4} alkyl, C_{2-4} alkenyl, C_{3-6} cycloalkyl, C_{3-6} cycloalkyl C_{1-3} alkyl or (heterocyclic group) C_{1-3} alkyl; and wherein R^6 may be optionally substituted on carbon by one methoxy, ethoxy or trifluoromethyl;

R^3 is hydrogen;

R^4 is C_{3-6} cycloalkyl, C_{3-6} cycloalkyl C_{1-4} alkyl, heterocyclyl or 1-methoxyprop-2-yl;

R^5 is C_{1-6} alkyl;

or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof.

Therefore in another aspect of the invention, there is provided a compound of formula (I) (as depicted above) wherein:

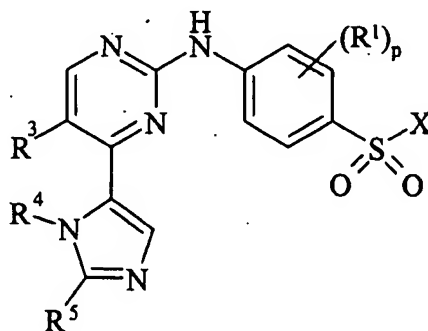
p is 0;

R^2 is $R^6-NH-R^6-NH_2$ - wherein R^6 is C_{1-4} alkyl, C_{2-4} alkenyl, C_{3-6} cycloalkyl, C_{3-6} cycloalkyl C_{1-3} alkyl or (heterocyclic group) C_{1-3} alkyl; and wherein R^6 may be optionally substituted on carbon by one methoxy, ethoxy or trifluoromethyl;

Please amend specification passage at page 10, lines 6-11 as follows:

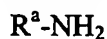
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Process c) for compounds of formula (I) where R^2 is amino or a group $R^6-NH-R^6-NH_2$;
reacting a pyrimidine of formula (VI):



(VI)

wherein X is a displaceable group; with an amine of formula (VII):



(VII)

wherein R^a is hydrogen or R^6 ;